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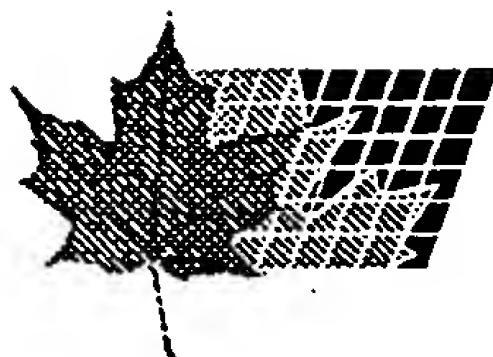
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(54) VOIE DE SYNTHÈSE BIOLOGIQUE DES GENES DES 1-  
DESOXY-D-XYLULOSE

(54) GENES OF THE 1-DESOXY-D-XYLULOSE BIOSYNTHETIC  
PATHWAY

(57) The invention relates to the 1-desoxy- D-xylulose- 5-phosphate reductoisomerase gene, the 1-desoxy- D-xylulose- 5-phosphate- synthase gene and the gcpE gene of the 1-desoxy- D-xylulose biosynthetic pathway and to their use for transforming vectors, host organisms and plants and for determining substances that inhibit this biosynthetic pathway.



**PCT**  
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(21) Internationales Aktenzeichen: PCT/EP99/07055 (22) Internationales Anmeldedatum: 22. September 1999 (22.09.99)  (30) Prioritätsdaten: 198 43 279.8      22. September 1998 (22.09.98)    DE 199 23 567.8      21. Mai 1999 (21.05.99)      DE  (71)(72) Anmelder und Erfinder: JOMAA, Hassan [DE/DE]; Breslauer Strasse 24, D-35398 Gießen (DE).  (74) Anwälte: PANTEN, Kirsten usw.; Reichel und Reichel, Park- strasse 13, D-60322 Frankfurt am Main (DE).		(81) Bestimmungsstaaten: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO Patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), eurasisches Patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), europäisches Patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI Patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  Veröffentlicht <i>Ohne internationalen Recherchenbericht und erneut zu veröffentlichen nach Erhalt des Berichts.</i>	
(54) Title: GENES OF THE 1-DESOXY-D-XYLULOSE BIOSYNTHETIC PATHWAY (54) Bezeichnung: GENE DES 1-DESOXY-D-XYLULOSE-BIOSYNTHESEWEGS (57) Abstract <p>The invention relates to the 1-desoxy- D-xylulose- 5-phosphate reductoisomerase gene, the 1-desoxy- D-xylulose- 5-phosphate-synthase gene and the gcpE gene of the 1-desoxy- D-xylulose biosynthetic pathway and to their use for transforming vectors, host organisms and plants and for determining substances that inhibit this biosynthetic pathway.</p> (57) Zusammenfassung <p>Die vorliegende Erfindung betrifft das 1-Desoxy- D-xylulose- 5-phosphatreduktisomerase -Gen, das 1-Desoxy- D-xylulose- 5-phosphat- Synthase- Gen und das gcpE-Gen des 1-Desoxy- D-xylulose- Biosynthesewegs und ihre Verwendung zur Transformation von Vektoren, Wirtsorganismen und Pflanzen und zur Bestimmung von Stoffen, die diesen Biosyntheseweg inhibieren.</p>			

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Claims

1. DNA sequences which code for a polypeptide with the amino acid sequence shown in SEQ ID no. 2 or for an analogue or derivative of the polypeptide according to SEQ ID no. 2, in which one or more amino acids have been deleted, added or replaced by other amino acids, wherein the enzymatic action of the polypeptide is retained, and which sequences originate from parasites, wherein sequence variations occurring within the framework of natural strain variability are included.
2. DNA sequences which code for a polypeptide with the amino acid sequence shown in SEQ ID no. 4 or for an analogue or derivative of the polypeptide according to SEQ ID no. 4, in which one or more amino acids have been deleted, added or replaced by other amino acids, wherein the enzymatic action of the polypeptide is retained, and which sequences originate from parasites, wherein sequence variations occurring within the framework of natural strain variability are included.
3. DNA sequences which code for a polypeptide with the amino acid sequence shown in SEQ ID no. 6 or for an analogue or derivative of the polypeptide according to SEQ ID no. 6, in which one or more amino acids have been deleted, added or replaced by other amino acids wherein the catalytic function of the polypeptide is retained.

- 8 -

4. DNA sequence according to one of claims 1 to 3, characterised in that it also comprises functional regulation signals, in particular promoters, operators, enhancers, ribosomal binding sites.
- 5
5. DNA sequence with the following sub-sequences
- 10 i) promoter which is active in viruses, eukaryotes and prokaryotes and ensures the formation of an RNA in the intended target tissue or target cells,
- ii) DNA sequences according to one of claims 1 to 3,
- 15 iii) 3' untranslated sequence which, in viruses, eukaryotes and prokaryotes, results in the addition of poly(A) residues onto the 3' end of the RNA.
6. Process for the production of transgenic viruses, eukaryotes and prokaryotes for modifying the
- 20 isoprenoid content, characterised in that a DNA sequence according to claim 4 or 5 is transferred and incorporated into the genome of viruses, eukaryotic and prokaryotic cells with or without use of a vector.
- 25
7. Transgenic systems, in particular plants and plant cells which contain one or more DNA sequences according to claims 1 to 5 as "foreign" or "additional" DNA, which sequences are expressed.
- 30
8. Expression vector containing one or more DNA sequences according to claims 1 to 5.

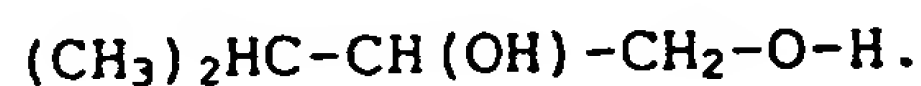
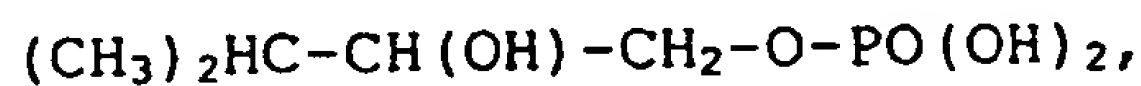
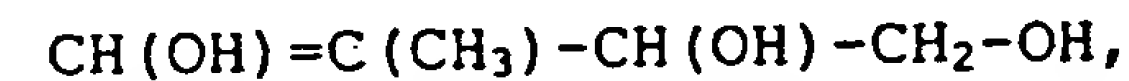
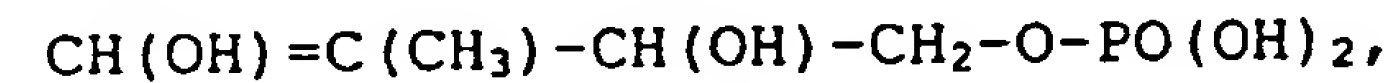
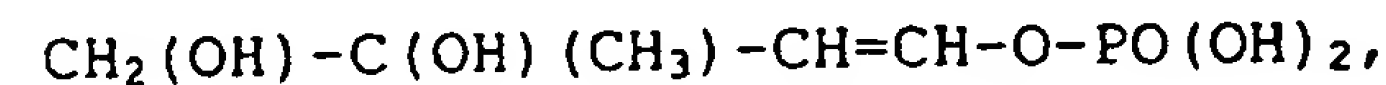
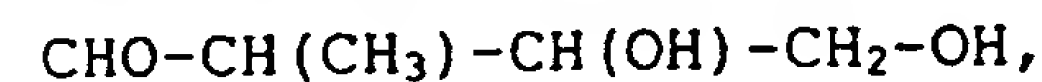
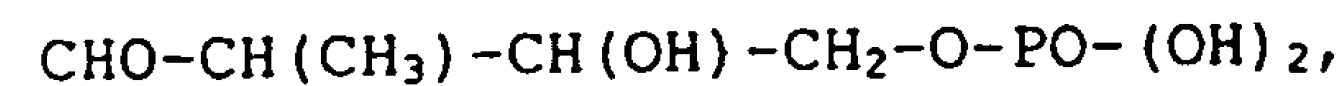
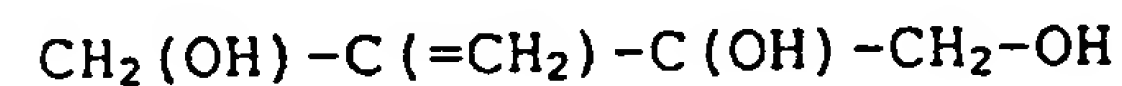
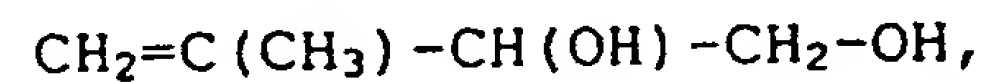
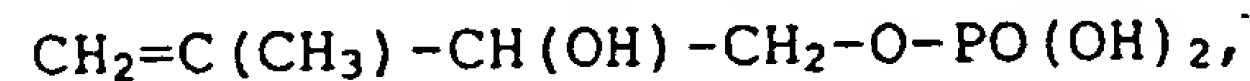
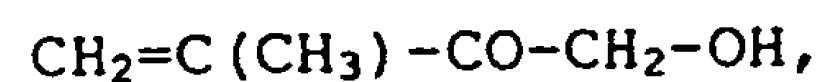
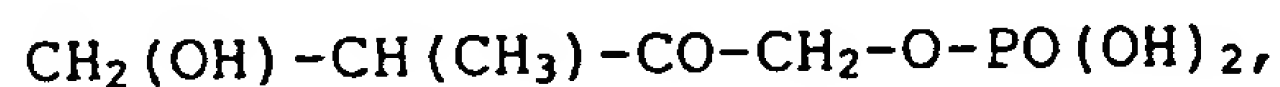
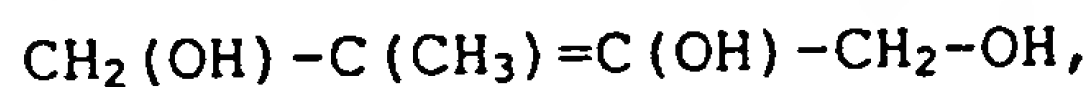
- 18 -

9. Protein which is involved in the 1-deoxy-D-xylulose 5-phosphate metabolic pathway and a) is coded by DNA sequences SEQ ID no. 1, 3 or 5 or b) is coded by DNA sequences which hybridise with DNA sequences SEQ ID no. 1, 3, 5 or fragments of these DNA sequences in the DNA region which codes for the mature protein.
10. Protein according to claim 9, obtainable from the culture supernatants of parasites or from the disrupted parasites and purification by chromatographic and electrophoretic methods.
11. Protein according to one of claims 9 and 10, characterised in that it a) is the product of viral, prokaryotic or eukaryotic expression of exogenous DNA, b) is coded by sequences SEQ ID no. 1, 3 or 5 or is coded by DNA sequences which hybridise with DNA sequences SEQ ID no. 1, 3, 5 or fragments of these DNA sequences in the DNA region which codes for the mature protein, or c) is coded by DNA sequences which would hybridise without degeneration of the genetic code with the sequences defined in b) and which code for a polypeptide with a corresponding amino acid sequence.
12. Protein according to one of the preceding claims, characterised in that it comprises the amino acid sequences SEQ ID no. 2, 4 or 6.
13. Process for determining the enzymatic activity of the gcpE protein, characterised in that phosphorylation of a sugar or of a phosphorus sugar or of a precursor of isoprenoid biosynthesis, in

- 9 -

particular the phosphorylation of 2-C-methyl-D-erythritol, 2-C-methyl-D-erytritol phosphate, in particular 2-C-methyl-D-erythritol 4-phosphate, 2-C-methyl-D-erythrose, 2-C-methyl-D-erythrose phosphate, in particular 2-C-methyl-D-erythrose 4-phosphate, and of phosphate and alcohol precursors, is detected.

14. Process according to claim 13, characterised in that phosphorylation of the following phosphates or alcohols is detected:



- 1 -

15. Process for the combined determination of the enzymatic activity of DOXP synthase and of DOXP reductase, characterised in that the conversion of glyceraldehyde 3-phosphate into 2-C-methylerythritol 4-phosphate is detected.
16. Process for screening a compound for the treatment of infectious processes in humans and animals, wherein the process comprises:
- a) provision of a host cell which contains a recombinant expression vector, wherein the vector comprises at least a portion of the oligonucleotide sequence according to SEQ ID no. 1, SEQ ID no. 3 or SEQ ID no. 5 or variants or analogues thereof, and moreover of a compound suspected to have antimycotic, antibiotic, antiparasitic or antiviral action in humans and animals,
  - b) bringing the host cell into contact with the compound and
  - c) determining the antimicrobial, antimycotic, antibiotic, antiparasitic or antiviral action of the compound.
17. Process for screening for compounds for treating plants, wherein the process comprises:
- a) provision of a host cell which contains a recombinant expression vector, wherein the vector comprises at least a portion of the oligonucleotide sequence according to SEQ ID no. 1, SEQ ID no. 3 or SEQ ID no. 5 or variants or analogues thereof, and moreover of a compound suspected to have antimicrobial,



- 9 -

antiviral, antiparasitic, bactericidal,  
fungicidal or herbicidal action in plants,

b) bringing the host cell into contact with the  
compound and

5 c) determining the antimicrobial, antiviral,  
antiparasitic, bactericidal, fungicidal or  
herbicidal action of the compound.

10 18. Use of DNA according to one of claims 1 to 5 or of  
proteins according to one of claims 9 to 12 or of  
transgenic systems according to claim 7 for the  
prevention or treatment of diseases in humans and  
animals.

- 1 -

Genes of the 1-deoxy-D-xylulose biosynthesis pathway

5 The present invention relates to DNA sequences which, when incorporated into the genome of viruses, eukaryotes and prokaryotes, modify isoprenoid biosynthesis and to a genetic engineering process for the production of these transgenic viruses, eukaryotes and prokaryotes. The invention also relates to a process for the identification of substances having herbicidal, antimicrobial, antiparasitic, antiviral, fungicidal, bactericidal action in plants and antimicrobial, antiparasitic, antimycotic, antibacterial and antiviral action in humans and animals.

15 The biosynthesis pathway for the formation of isoprenoids via the classical acetate/mevalonate pathway and an alternative mevalonate-independent biosynthesis pathway, the deoxy-D-xylulose phosphate pathway is already known (Rohmer, M., Knani, M., Simonin, P., Sutter, B. and Sahm, H. (1993): *Biochem. J.* 295: 517-524).

25 It is, however, not known how and by which pathways it is possible to bring about a change in the isoprenoid concentration in viruses, eukaryotes and prokaryotes by means of the deoxy-D-xylulose phosphate pathway. Figure 1 shows this biosynthesis pathway.

30 DNA sequences are consequently provided which code for 1-deoxy-D-xylulase 5-phosphate synthase (DOXP synthase), 1-deoxy-D-xylulose 5-phosphate reductoisomerase (DOXP reductoisomerase) or the gcpE protein. All three genes and enzymes are involved in isoprenoid biosynthesis.

-2-

(Translator's comment: The portion at the beginning of the next paragraph enclosed in square brackets corresponds to the beginning of the sentence which finishes on page 2, line 1 of the original).

[The gcpE protein has a kinase function and catalyses the phosphorylation of a sugar or a phosphorus sugar or a precursor of isoprenoid biosynthesis, in particular the phosphorylation of 2-C-methyl-D-erythritol, 2-C-methyl-D-erythritol phosphate, in particular 2-C-methyl-D-erythritol 4-phosphate, 2-C-methyl-D-erythrose, 2-C-methyl-D-erythrose] phosphate, in particular 2-C-methyl-D-erythrose 4-phosphate. In the precursor of isoprenoid synthesis, the gcpE protein in particular catalyses the phosphorylation of the following substances:

CH<sub>2</sub>(OH)-C(CH<sub>3</sub>)=C(OH)-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 15 CH<sub>2</sub>(OH)-C(CH<sub>3</sub>)=C(OH)-CH<sub>2</sub>-OH,  
 CH<sub>2</sub>(OH)-CH(CH<sub>3</sub>)-CO-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 CH<sub>2</sub>(OH)-CH(CH<sub>3</sub>)-CO-CH<sub>2</sub>OH  
 CH<sub>2</sub>=C(CH<sub>3</sub>)-CO-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 CH<sub>2</sub>=C(CH<sub>3</sub>)-CO-CH<sub>2</sub>-OH,  
 20 CH<sub>2</sub>=C(CH<sub>3</sub>)-CH(OH)-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 CH<sub>2</sub>=C(CH<sub>3</sub>)-CH(OH)-CH<sub>2</sub>-OH,  
 CH<sub>2</sub>(OH)-C(=CH<sub>2</sub>)-C(OH)-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 CH<sub>2</sub>(OH)-C(=CH<sub>2</sub>)-C(OH)-CH<sub>2</sub>-OH  
 CHO-CH(CH<sub>3</sub>)-CH(OH)-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 25 CHO-CH(CH<sub>3</sub>)-CH(OH)-CH<sub>2</sub>-OH,  
 CH<sub>2</sub>(OH)-C(OH)(CH<sub>3</sub>)-CH=CH-O-PO(OH)<sub>2</sub>,  
 CH<sub>2</sub>(OH)-C(OH)(CH<sub>3</sub>)-CH=CH-OH  
 CH(OH)=C(CH<sub>3</sub>)-CH(OH)-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 CH(OH)=C(CH<sub>3</sub>)-CH(OH)-CH<sub>2</sub>-OH,  
 30 (CH<sub>3</sub>)<sub>2</sub>HC-CO-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 (CH<sub>3</sub>)<sub>2</sub>HC-CO-CH<sub>2</sub>-O-H,  
 (CH<sub>3</sub>)<sub>2</sub>HC-CH(OH)-CH<sub>2</sub>-O-PO(OH)<sub>2</sub>,  
 (CH<sub>3</sub>)<sub>2</sub>HC-CH(OH)-CH<sub>2</sub>-O-H.

- 2 -

DOXP synthase catalyses the condensation of pyruvate and  
glyceraldehyde 3-phosphate to yield 1-deoxy-D-xylulose  
5-phosphate and DOXP reductoisomerase catalyses the  
5 conversion of 1-deoxy-D-xylulose 5-phosphate into  
2-C-methyl-D-erythritol 4-phosphate (c.f. Fig. 1).

The invention relates to the following DNA sequences:  
DNA sequences which code for a polypeptide with the amino  
10 acid sequence shown in SEQ ID no. 2 or for an analogue or  
derivative of the polypeptide according to SEQ ID no. 2,  
in which one or more amino acids have been deleted, added  
or replaced by other amino acids, wherein the enzymatic  
action of the polypeptide is retained, and which  
15 sequences originate from parasites, wherein sequence  
variations occurring within the framework of natural  
strain variability are included,

DNA sequences which code for a polypeptide with the amino  
20 acid sequence shown in SEQ ID no. 4 or for an analogue or  
derivative of the polypeptide according to SEQ ID no. 4,  
in which one or more amino acids have been deleted, added  
or replaced by other amino acids, wherein the enzymatic  
action of the polypeptide is retained, and which  
25 sequences originate from parasites, wherein sequence  
variations occurring within the framework of natural  
strain variability are included,

and DNA sequences which code for a polypeptide with the  
30 amino acid sequence shown in SEQ ID no. 6 or for an  
analogue or derivative of the polypeptide according to  
SEQ ID no. 6, in which one or more amino acids have been

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*Amendments*

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- 3 -

deleted, added or replaced by other amino acids, wherein the catalytic function of the polypeptide is retained.

25     The genes and the gene products thereof (polypeptides)  
are shown with their primary structure and are assigned  
as follows:

SEQ ID no. 1: 1-deoxy-D-xylulose 5-phosphate reducto-  
isomerase gene

30     SEQ ID no. 2: 1-deoxy-D-xylulose 5-phosphate reducto-  
isomerase

SEQ ID no. 3: 1-deoxy-D-xylulose 5-phosphate synthase  
gene

SEQ ID no. 4: 1-deoxy-D-xylulose 5-phosphate synthase

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- 4 -

SEQ ID no. 5: gcpE gene

SEQ ID no. 6: gcpE proteins.

5 The DNA sequences all originate from *Plasmodium falciparum*.

10 Apart from the DNA sequences stated in the sequence listing, suitable sequences are also those which, as a result of the degeneration of the genetic code, have another DNA sequence, but code for the same peptide or for an analogue or derivative of the polypeptide, in which one or more amino acids have been deleted, added or replaced by other amino acids.

15 The sequences according to the invention are suitable for the expression of genes in viruses, eukaryotes and prokaryotes which are responsible for isoprenoid biosynthesis in the 1-deoxy-D-xylulose pathway.

20 According to the invention, eukaryotes or eukaryotic cells include animal cells, plant cells, algae, yeasts, fungi, while prokaryotes or prokaryotic cells include bacteria, archaebacteria and eubacteria.

25 When a DNA sequence is incorporated into a genome on which the above-stated DNA sequence is located, expression of the above-described genes in viruses, eukaryotes and prokaryotes is enabled. The viruses, eukaryotes and prokaryotes transformed according to the  
30 invention are cultivated in a manner known per se and the isoprenoid formed during such cultivation is isolated and optionally purified. Not all isoprenoids need to be

- 5 -

isolated as in some case the isoprenoids are released directly into the ambient air.

5 The invention furthermore relates to a process for the production of transgenic viruses, eukaryotes and prokaryotes in order to modify the isoprenoid content, which process comprises the following steps.

- 10 a) Production of a DNA sequence with the following sub-sequences
- i) promoter which is active in viruses, eukaryotes and prokaryotes and ensures the formation of an RNA in the intended target tissue or target cells,
  - 15 ii) DNA sequence which codes for a polypeptide with the amino acid sequence shown in SEQ ID no. 2, 4 or 6 or for an analogue or derivative of the polypeptide according to SEQ ID no. 2, 4 or 6,
  - 20 iii) 5' and 3' untranslated sequence which enables or enhances expression of the stated genes in viruses, eukaryotes and prokaryotes,
- b) transfer and incorporation of the DNA sequence into the genome of viruses, prokaryotic or eukaryotic cells with or without the use of a vector (for
- 25 example plasmid, viral DNA).

The intact, whole plants may be regenerated from plant cells transformed in this manner.

30 The protein-coding sequences with the nucleotide sequences SEQ ID no. 1, SEQ ID no. 3 and SEQ ID no. 5 may be provided with a promoter which ensures transcription in certain organs or cells, which promoter is coupled in

- 6 -

sense orientation (3' end of the promoter to the 5' end of the coding sequence) to the sequence which codes the protein to be formed. A termination signal which determines termination of mRNA synthesis is attached to the 3' end of the coding sequence. In order to direct the protein which is to be expressed to certain subcellular compartments, such as chloroplasts, amyloplasts, mitochondria, vacuoles, cytosol or intercellular spaces, a further sequence which codes for a so-called signal sequence or a transit peptide may be inserted between the promoter and the coding sequence. In some cases, it is necessary to insert sequences which code for a signal at the COOH terminus of the protein. The sequence must be in the same reading frame as the coding sequence of the protein. A large number of cloning vectors is available in order to prepare for the introduction of the DNA sequences according to the invention into higher plants, which vectors contain a replication signal for *E. coli* and a marker which permits selection of the transformed cells. Depending upon the method by which desired genes are introduced into the plant, further DNA sequences may be required. If, for example, the Ti or Ri plasmid is used to transform the plant cells, at least one right border, but frequently the right border and left border of the Ti and Ri plasmid T-DNA must be inserted as a flanking region into the genes to be introduced. The use of T-DNA for transforming plant cells has been intensively investigated and comprehensively described in EP 120516; Hoekama in "The Binary Plant Vector System", Offset-drukkerij Kanters B.V. Alblasserdam (1985), chapter V; Fraley et al., *Crit.Rev.Plant Sci.* 4, 1-46 and An et al. (1985) *EMBO J.* 4, 277-287. Once the introduced DNA has been incorporated into the genome, it is



generally stable and is also retained in the descendants of the originally transformed cells. It normally contains a selection marker, which imparts to the transformed plant cells resistance to a biocide or an antibiotic, such as kanamycin, G 418, bleomycin, hygromycin or phosphinotricin and others. The particular marker used is thus intended to allow selection of transformed cells from cells lacking the inserted DNA.

Many techniques are available for introducing DNA into a plant. These techniques include transformation with the assistance of agrobacteria, for example *Agrobacterium tumefaciens*, protoplast fusion, microinjection of DNA, electroporation, as well as ballistic methods and virus infection. Whole plants may then be regenerated from the transformed plant material in a suitable medium which may contain antibiotics or biocides for selection purposes. No particular requirements are placed upon the plasmids for injection and electroporation. However, if whole plants are to be regenerated from such transformed cells, a selectable marker gene must be present. The transformed cells grow in the plants in the conventional manner (McCormick et al. (1986), *Plant Cell Reports* 5, 81-84). The plants may be cultivated normally and be crossed with plants which have the same transformed genome or other genomes. The resultant individuals have the corresponding phenotypic properties.

The present invention also provides expression vectors which contain one or more of the DNA sequences according to the invention. Such expression vectors are obtained by providing the DNA sequences according to the invention with suitable functional regulation signals. Such

- 8 -

regulation signals are DNA sequences which are responsible for expression, for example promoters, operators, enhancers, ribosomal binding sites, and are recognised by the host organism.

5

Further regulation signals, which for example control replication or recombination of the recombinant DNA in the host organism, may optionally also be a constituent part of the expression vector.

10

The host organisms transformed with the DNA sequences or expression vectors according to the invention are also provided by the present invention.

15

Suitable host cells and organisms for expressing the enzymes according to the invention are those which comprise no intrinsic enzymes with the function of DOXP synthase, DOXP reductoisomerase or the gcpE protein. This is the case for archaebacteria, animals, fungi, slime

20

moulds and some eubacteria. The absence of such intrinsic enzyme activity substantially facilitates detection and purification of the recombinant enzymes. As a

25

consequence, it is also for the first time possible straightforwardly to measure, in crude extracts from the host cells, the activity and in particular the inhibition of the activity of the recombinant enzymes according to the invention by various chemicals and pharmaceuticals.

30

The enzymes according to the invention are advantageously then expressed in eukaryotic cells if post-translational modification and native folding of the polypeptide chain is to be achieved. Moreover, depending upon the expression system, it is ensured when expressing genomic

- 9 -

DNA sequences that introns are eliminated by splicing the DNA and the enzymes are produced in the polypeptide sequences characteristic to the parasites. Using recombinant DNA techniques, sequences coding for introns may be eliminated from or inserted for experimental purposes into the DNA sequences to be expressed.

The protein may be isolated from the host cell or the culture supernatant of the host cell using methods known to the person skilled in the art. In vitro reactivation of the enzymes may also be required.

In order to facilitate purification, the enzymes according to the invention or sub-sequences of the enzymes may be expressed as fusion proteins with various peptide chains. Oligo-histidine sequences and sequences derived from glutathione S-transferase, thioredoxin or calmodulin-binding peptides are particularly suitable for this purpose.

20

The enzymes according to the invention or sub-sequences of the enzymes may furthermore be expressed as fusion proteins with such peptide chains known to the person skilled in the art that the recombinant enzymes are transported into the extracellular medium or into certain compartments of the host cells. Both purification and investigation of the biological activity of the enzymes may consequently be facilitated.

When expressing the enzymes according to the invention, it may prove convenient to modify individual codons. Purposeful replacement of bases in the coding region may here also be advisable if the codons used in the

- 10 -

parasites differ from the codon use in the heterologous expression system, in order to ensure optimal synthesis of the protein.

5 The enzymes according to the invention may furthermore be obtained under standardised conditions by *in vitro* translation by methods known to the person skilled in the art. Systems suitable for this purpose are rabbit reticulocyte and wheat germ extracts and bacterial  
10 lysates. *In vitro* transcribed mRNA may also be translated into *Xenopus* oocytes.

Oligo- and polypeptides, the sequences of which are derived from the peptide sequence of the enzymes  
15 according to the invention, may be obtained by chemical synthesis. Given appropriate selection of the sequences, such peptides have properties which are characteristic of the enzymes according to the invention. Such peptides may be produced in large quantities and are particularly  
20 suitable for investigating the kinetics of enzyme activity, regulation of enzyme activity, the three-dimensional structure of the enzymes, inhibition of enzyme activity by various chemicals and pharmaceuticals and the binding geometry and binding affinity of various  
25 ligands.

DNA with the nucleotides from sequences SEQ ID no. 1, 3 and 5 are preferably used for the recombinant production of the enzymes according to the invention.

30

The invention accordingly moreover relates to a process for screening for compounds which inhibit the deoxy-D-xylulose phosphate metabolic pathway. According to this

- 11 -

process, a host organism, which contains a recombinant expression vector, wherein the vector comprises at least a portion of the oligonucleotide sequence according to SEQ ID no. 1, SEQ ID no. 3 or SEQ ID no. 5 or variants or homologues thereof, is provided, as is a compound which is suspected to have antimicrobial, antiparasitic, antibacterial, antiviral and antimycotic action in humans and animals or an antimicrobial, antiviral, bactericidal, herbicidal or fungicidal activity in plants. The host organism is then brought into contact with the compound and the activity of the compound determined.

The present invention also provides methods for determining the enzymatic activity of the gcpE protein. Said activity may be determined using known methods. Determination is performed by detecting the phosphorylation of a sugar or of a phosphorus sugar or of a precursor of isoprenoid biosynthesis, in particular the phosphorylation of 2-C-methyl-D-erythritol, 2-C-methyl-D-erythritol phosphate, in particular 2-C-methyl-D-erythritol 4-phosphate, 2-C-methyl-D-erythrose, 2-C-methyl-D-erythrose phosphate, in particular 2-C-methyl-D-erythrose 4-phosphate. The present invention also provides the use of this measurement method for identifying substances which inhibit the activity of the particular enzymes.

The enzymatic activity of DOXP synthase and DOXP reductoisomerase may be detected in a single step by determining the conversion of glyceraldehyde 3-phosphate into 2-C-methylerythritol 4-phosphate.

- 12 -

Determination of the activities of DOXP synthase and DOXP reductoisomerase proceeds analogously. Fluorimetric methods described by Querol et al. are also suitable for determining DOXP synthase activity (Querol et al.,  
5 abstracts, 4<sup>th</sup> European Symposium on Plant Isoprenoids, Barcelona, 21-23 April 1999).

WO 00/17233

PCT/EP99/07055

- 1 -

## SEQUENCE LISTING

&lt;110&gt; Jomaa, Hassan

&lt;120&gt; Genes of the 1-deoxy-D-xylulose biosynthesis pathway

&lt;130&gt; 15696

&lt;140&gt; PCT/EP99

&lt;141&gt; 1999-09-22

&lt;150&gt; DE19923567.8

&lt;151&gt; 1999-05-22

&lt;150&gt; DE19843279.8

&lt;151&gt; 1998-09-22

&lt;160&gt; 6

&lt;170&gt; PatentIn Ver. 2.1

&lt;210&gt; 1

&lt;211&gt; 1467

&lt;212&gt; DNA

&lt;213&gt; Plasmodium falciparum

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (1)..(1467)

&lt;220&gt;

&lt;221&gt; gene

&lt;222&gt; (1)..(1467)

&lt;220&gt;

&lt;221&gt; mRNA

&lt;222&gt; (1)..(1467)

&lt;400&gt; 1

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1				5				10						15		

aat	gat	tta	gta	ata	aat	aat	aca	tca	aaa	tgt	gtt	tcc	att	gaa	aga	96
Asn	Asp	Leu	Val	Ile	Asn	Asn	Thr	Ser	Lys	Cys	Val	Ser	Ile	Glu	Arg	
		20					25						30			

aga	aaa	aat	aac	gca	tat	ata	aat	tat	ggt	ata	gga	tat	aat	gga	cca	144
Arg	Lys	Asn	Asn	Ala	Tyr	Ile	Asn	Tyr	Gly	Ile	Gly	Tyr	Asn	Gly	Pro	
		35					40					45				

gat	aat	aaa	ata	aca	aag	agt	aga	aga	tgt	aaa	aga	ata	aag	tta	tgc	192
Asp	Asn	Lys	Ile	Thr	Lys	Ser	Arg	Arg	Cys	Lys	Arg	Ile	Lys	Leu	Cys	
		50				55					60					



WO 00/17233

PCT/EP99/07055

- 2 -

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Lys Lys Asp Leu Ile Asp Ile Gly Ala Ile Lys Lys Pro Ile Asn Val	
65 70 75 80	
gca att ttt gga agt act ggt agt ata ggt acg aat gct tta aat ata	288
Ala Ile Phe Gly Ser Thr Gly Ser Ile Gly Thr Asn Ala Leu Asn Ile	
85 90 95	
ata agg gag tgt aat aaa att gaa aat gtt ttt aat gtt aaa gca ttg	336
Ile Arg Glu Cys Asn Lys Ile Glu Asn Val Phe Asn Val Lys Ala Leu	
100 105 110	
tat gtg aat aag agt gtg aat gaa tta tat gaa caa gct aga gaa ttt	384
Tyr Val Asn Lys Ser Val Asn Glu Leu Tyr Glu Gln Ala Arg Glu Phe	
115 120 125	
tta cca gaa tat ttg tgt ata cat gat aaa agt gta tat gaa gaa tta	432
Leu Pro Glu Tyr Leu Cys Ile His Asp Lys Ser Val Tyr Glu Glu Leu	
130 135 140	
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Lys Glu Leu Val Lys Asn Ile Lys Asp Tyr Lys Pro Ile Ile Leu Cys	
145 150 155 160	
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Gly Asp Glu Gly Met Lys Glu Ile Cys Ser Ser Asn Ser Ile Asp Lys	
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Ile Val Ile Gly Ile Asp Ser Phe Gln Gly Leu Tyr Ser Thr Met Tyr	
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Ala Ile Met Asn Asn Lys Ile Val Ala Leu Ala Asn Lys Glu Ser Ile	
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210 215 220	
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Leu Asp Asn Asn Lys Val Leu Lys Thr Lys Cys Leu Gln Asp Asn Phe	
245 250 255	
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Ser Lys Ile Asn Asn Ile Asn Lys Ile Phe Leu Cys Ser Ser Gly Gly	
260 265 270	
cca ttt caa aat tta act atg gac gaa tta aaa aat gta aca tca gaa	864
Pro Phe Gln Asn Leu Thr Met Asp Glu Leu Lys Asn Val Thr Ser Glu	
275 280 285	



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 305 310 315 320

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 325 330 335

gaa tgc att ata cat tct tgt gtt gaa ttt ata gac aaa tca gta ata 1056  
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 340 345 350

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 355 360 365

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 370 375 380

cag gtt tca act ctt aca ttt cat aaa cct tct tta gaa cat ttc ccg 1200  
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 385 390 395 400

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 405 410 415

act gta cta aat gcg tca aat gaa ata gct aac aac tta ttt ttg aat 1296  
 Thr Val Leu Asn Ala Ser Asn Glu Ile Ala Asn Asn Leu Phe Leu Asn  
 420 425 430

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 435 440 445

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&lt;210&gt; 2

&lt;211&gt; 488

&lt;212&gt; PRT

&lt;213&gt; Plasmodium falciparum

WO 00/17233

PCT/EP99/07055

- 4 -

&lt;400&gt; 2

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 Arg Lys Asn Asn Ala Tyr Ile Asn Tyr Gly Ile Gly Tyr Asn Gly Pro  
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 Asp Asn Lys Ile Thr Lys Ser Arg Arg Cys Lys Arg Ile Lys Leu Cys  
 50 55 60  
 Lys Lys Asp Leu Ile Asp Ile Gly Ala Ile Lys Lys Pro Ile Asn Val  
 65 70 75 80  
 Ala Ile Phe Gly Ser Thr Gly Ser Ile Gly Thr Asn Ala Leu Asn Ile  
 85 90 95  
 Ile Arg Glu Cys Asn Lys Ile Glu Asn Val Phe Asn Val Lys Ala Leu  
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 Tyr Val Asn Lys Ser Val Asn Glu Leu Tyr Glu Gln Ala Arg Glu Phe  
 115 120 125  
 Leu Pro Glu Tyr Leu Cys Ile His Asp Lys Ser Val Tyr Glu Glu Leu  
 130 135 140  
 Lys Glu Leu Val Lys Asn Ile Lys Asp Tyr Lys Pro Ile Ile Leu Cys  
 145 150 155 160  
 Gly Asp Glu Gly Met Lys Glu Ile Cys Ser Ser Asn Ser Ile Asp Lys  
 165 170 175  
 Ile Val Ile Gly Ile Asp Ser Phe Gln Gly Leu Tyr Ser Thr Met Tyr  
 180 185 190  
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 Val Ser Ala Gly Phe Phe Leu Lys Lys Leu Leu Asn Ile His Lys Asn  
 210 215 220  
 Ala Lys Ile Ile Pro Val Asp Ser Glu His Ser Ala Ile Phe Gln Cys  
 225 230 235 240  
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 245 250 255  
 Ser Lys Ile Asn Asn Ile Asn Lys Ile Phe Leu Cys Ser Ser Gly Gly  
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 Pro Phe Gln Asn Leu Thr Met Asp Glu Leu Lys Asn Val Thr Ser Glu  
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 290 295 300

WO 00/17233

PCT/EP99/07055

- 5 -

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 Phe Leu Phe Asp Val Asp Tyr Asn Asp Ile Glu Val Ile Val His Lys  
 325 330 335  
 Glu Cys Ile Ile His Ser Cys Val Glu Phe Ile Asp Lys Ser Val Ile  
 340 345 350  
 Ser Gln Met Tyr Tyr Pro Asp Met Gln Ile Pro Ile Leu Tyr Ser Leu  
 355 360 365  
 Thr Trp Pro Asp Arg Ile Lys Thr Asn Leu Lys Pro Leu Asp Leu Ala  
 370 375 380  
 Gln Val Ser Thr Leu Thr Phe His Lys Pro Ser Leu Glu His Phe Pro  
 385 390 395 400  
 Cys Ile Lys Leu Ala Tyr Gln Ala Gly Ile Lys Gly Asn Phe Tyr Pro  
 405 410 415  
 Thr Val Leu Asn Ala Ser Asn Glu Ile Ala Asn Asn Leu Phe Leu Asn  
 420 425 430  
 Asn Lys Ile Lys Tyr Phe Asp Ile Ser Ser Ile Ile Ser Gln Val Leu  
 435 440 445  
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&lt;213&gt; Plasmodium falciparum

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&lt;221&gt; gene

&lt;222&gt; (1)..(3870)

&lt;220&gt;

&lt;221&gt; mRNA

&lt;222&gt; (1)..(3870)

&lt;400&gt; 3

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WO 00/17233

PCT/EP99/07055

- 6 -

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Met Ile Phe Asn Tyr Val Phe Phe Lys Asn Phe Val Pro Val Val  
1 5 10 15  
cta tac att ctc ctt ata ata tat att aac tta aat ggc atg aat aat 218  
Leu Tyr Ile Leu Leu Ile Ile Tyr Ile Asn Leu Asn Gly Met Asn Asn  
20 25 30  
aaa aat caa ata aaa aca gaa aaa att tat ata aag aaa ttg aat agg 266  
Lys Asn Gln Ile Lys Thr Glu Lys Ile Tyr Ile Lys Lys Leu Asn Arg  
35 40 45  
ttg tca agg aaa aat tcg tta tgt agt tct aaa aat aaa ata gca tgc 314  
Leu Ser Arg Lys Asn Ser Leu Cys Ser Ser Lys Asn Lys Ile Ala Cys  
50 55 60  
ttg ttc gat ata gga aat gat gat aat aga aat acg aca tat ggc tat 362  
Leu Phe Asp Ile Gly Asn Asp Asp Asn Arg Asn Thr Thr Tyr Gly Tyr  
65 70 75  
aat gtg aat gtt aaa aat gat gat att aat tcc tta cta aaa aat aat 410  
Asn Val Asn Val Lys Asn Asp Asp Ile Asn Ser Leu Leu Lys Asn Asn  
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Tyr Ser Asn Lys Leu Tyr Met Asp Lys Arg Lys Asn Ile Asn Asn Val  
100 105 110  
att agt act aat aaa ata tct ggg tcc att tca aat att tgt agt aga 506  
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115 120 125  
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Asn Gln Lys Glu Asn Glu Gln Lys Arg Asn Lys Gln Arg Cys Leu Thr  
130 135 140  
caa tgt cac act tat aat atg tca cat gaa cag gac aaa cta gct aat 602  
Gln Cys His Thr Tyr Asn Met Ser His Glu Gln Asp Lys Leu Ala Asn  
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gat aat aat agg aat aat aaa aag aat ttt aat tta tta ttt ata aat 650  
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160 165 170 175  
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Tyr Phe Asn Leu Lys Arg Met Lys Asn Ser Leu Leu Asn Lys Asp Asn  
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195 200 205  
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210 215 220

WO 00/17233

PCT/EP99/07055

- 7 -

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aat gat aat aat gat tat aat aat aat aat agt tgt aat aat tta gga Asn Asp Asn Asn Asp Tyr Asn Asn Asn Asn Ser Cys Asn Asn Leu Gly 275 280 285	986
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tat ttc cca tta tta aaa tta att aat aat cca tca gat tta aaa aag Tyr Phe Pro Leu Leu Lys Leu Ile Asn Asn Pro Ser Asp Leu Lys Lys 420 425 430	1418
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Ser Ser Leu Glu Ile Gln Leu Leu Leu Tyr Ile Phe Asn Gln Pro	
465 470 475	
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Tyr Asp Asn Val Ile Tyr Asp Ile Gly His Gln Ala Tyr Val His Lys	
480 485 490 495	
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Ile Glu Ile Ser Asp Asn Ala Asn Val Thr Asn Asn Glu Arg Ile Phe	
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580 585 590	
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Asn Asn Tyr Ile Asn Pro Ser Asp Val Val Gly Arg Glu Asn Thr Asn	
595 600 605	
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Val Pro Asn Val Arg Asn Asp Asn His Asn Val Asp Lys Val His Ile	
610 615 620	
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625 630 635	
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Leu Asn Tyr Ile Ser Phe Leu Asn Ser Lys Ile Leu Ile Ile Tyr Asn	
640 645 650 655	
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660 665 670	

WO 00/17233

PCT/EP99/07055

- 9 -

aat aga cct ata ggt tct ata tca gat cat tta cat tat ttt gtt tct Asn Arg Pro Ile Gly Ser Ile Ser Asp His Leu His Tyr Phe Val Ser 675 680 685	2186
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gag aat aac att ttt gaa aat ttg aat tat gat tat att ggt gtt gtg Glu Asn Asn Ile Phe Glu Asn Leu Asn Tyr Asp Tyr Ile Gly Val Val 705 710 715	2282
aat ggt aat aat aca gaa gag ctc ttt aaa gta tta aat aat ata aaa Asn Gly Asn Asn Thr Glu Glu Leu Phe Lys Val Leu Asn Asn Ile Lys 720 725 730 735	2330
gaa aat aaa tta aaa aga gct act gtt ctt cat gta cgt aca aaa aaa Glu Asn Lys Leu Lys Arg Ala Thr Val Leu His Val Arg Thr Lys Lys 740 745 750	2378
tcg aat gat ttt ata aat tca aag agt cca ata agt ata ttg cac tct Ser Asn Asp Phe Ile Asn Ser Lys Ser Pro Ile Ser Ile Leu His Ser 755 760 765	2426
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tct aca aag tat gat gta aat aat aag aat aat aaa aat aat gat aat Ser Thr Lys Tyr Asp Val Asn Asn Lys Asn Asn Lys Asn Asn Asp Asn 800 805 810 815	2570
agt gaa att ata aaa tat gaa gat atg ttt tca aaa gag acg ttc aca Ser Glu Ile Ile Lys Tyr Glu Asp Met Phe Ser Lys Glu Thr Phe Thr 820 825 830	2618
gat ata tat aca aat gaa atg tta aaa tat tta aag aaa gat aga aat Asp Ile Tyr Thr Asn Glu Met Leu Lys Tyr Leu Lys Lys Asp Arg Asn 835 840 845	2666
ata ata ttc cta tct ccc gct atg tta gga gga tca gga ttg gtt aaa Ile Ile Phe Leu Ser Pro Ala Met Leu Gly Gly Ser Gly Leu Val Lys 850 855 860	2714
att agt gag cgt tat cca aat aat gta tat gat gta ggt ata gca gaa Ile Ser Glu Arg Tyr Pro Asn Asn Val Tyr Asp Val Gly Ile Ala Glu 865 870 875	2762
caa cat tct gta act ttc gca gca gct atg gca atg aat aag aaa tta Gln His Ser Val Thr Phe Ala Ala Ala Met Ala Met Asn Lys Lys Leu 880 885 890 895	2810



WO 00/17233

PCT/EP99/07055

- 10 -

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 900 905 910

caa att ata cat gat ctt aat tta caa aat ata cct tta aag gtt ata 2906  
 Gln Ile Ile His Asp Leu Asn Leu Gln Asn Ile Pro Leu Lys Val Ile  
 915 920 925

att gga aga agt gga tta gta gga gag gat ggg gca aca cat caa ggt 2954  
 Ile Gly Arg Ser Gly Leu Val Gly Glu Asp Gly Ala Thr His Gln Gly  
 930 935 940

ata tat gat tta tct tat ctt ggg aca ctt aac aat gca tat ata ata 3002  
 Ile Tyr Asp Leu Ser Tyr Leu Gly Thr Leu Asn Asn Ala Tyr Ile Ile  
 945 950 955

tct cca agt aat caa gtt gat ttg aaa aga gct ctt agg ttt gct tat 3050  
 Ser Pro Ser Asn Gln Val Asp Leu Lys Arg Ala Leu Arg Phe Ala Tyr  
 960 965 970 975

tta gat aag gac cat tct gtg tat ata cgt ata ccc aga atg aac ata 3098  
 Leu Asp Lys Asp His Ser Val Tyr Ile Arg Ile Pro Arg Met Asn Ile  
 980 985 990

tta agt gat aag tac atg aaa gga tat ttg aac att cat atg aaa aat 3146  
 Leu Ser Asp Lys Tyr Met Lys Gly Tyr Leu Asn Ile His Met Lys Asn  
 995 1000 1005

gag agc aaa aat atc gat gta aac gtg gat ata aac gat gat gta gat 3194  
 Glu Ser Lys Asn Ile Asp Val Asn Val Asp Ile Asn Asp Asp Val Asp  
 1010 1015 1020

aaa tat agt gaa gaa tat atg gac gat gat aat ttt ata aaa tcg ttt 3242  
 Lys Tyr Ser Glu Glu Tyr Met Asp Asp Asp Asn Phe Ile Lys Ser Phe  
 1025 1030 1035

att gga aaa tct aga att att aaa atg gat aat gaa aat aat aat aca 3290  
 Ile Gly Lys Ser Arg Ile Ile Lys Met Asp Asn Glu Asn Asn Asn Thr  
 1040 1045 1050 1055

aat gaa cat tat tca agc aga gga gat aca cag aca aaa aaa aaa aaa 3336  
 Asn Glu His Tyr Ser Ser Arg Gly Asp Thr Gln Thr Lys Lys Lys Lys  
 1060 1065 1070

gtt tgt atc ttt aac atg ggt agt atg ctt ttt aat gta att aat gct 3366  
 Val Cys Ile Phe Asn Met Gly Ser Met Leu Phe Asn Val Ile Asn Ala  
 1075 1080 1085

ata aaa gaa att gaa aaa gaa caa tat att tca cat aat tat tct ttt 3434  
 Ile Lys Glu Ile Glu Lys Glu Gln Tyr Ile Ser His Asn Tyr Ser Phe  
 1090 1095 1100

tca att gtt gat atg ata ttt tta aat cct tta gat aaa aat atg ata 3482  
 Ser Ile Val Asp Met Ile Phe Leu Asn Pro Leu Asp Lys Asn Met Ile  
 1105 1110 1115



WO 00/17233

PCT/EP99/07055

- 11 -

gat cat gta ata aaa caa aat aaa cat caa tat tta att act tat gaa 3530  
 Asp His Val Ile Lys Gln Asn Lys His Gln Tyr Leu Ile Thr Tyr Glu  
 1120 1125 1130 1135

gat aat act ata ggt ggt ttt tct aca cat ttc aat aat tat tta ata 3578  
 Asp Asn Thr Ile Gly Gly Phe Ser Thr His Phe Asn Asn Tyr Leu Ile  
 1140 1145 1150

gaa aat aat tat att aca aaa cat aac tta tat gtt cat aat att tat 3626  
 Glu Asn Asn Tyr Ile Thr Lys His Asn Leu Tyr Val His Asn Ile Tyr  
 1155 1160 1165

tta tct aat gag cca att gaa cat gca tct ttt aag gat caa caa gaa 3674  
 Leu Ser Asn Glu Pro Ile Glu His Ala Ser Phe Lys Asp Gln Gln Glu  
 1170 1175 1180

gtc gtc aaa atg gat aaa tgt agt ctt gtc aat aga att aaa aat tat 3722  
 Val Val Lys Met Asp Lys Cys Ser Leu Val Asn Arg Ile Lys Asn Tyr  
 1185 1190 1195

ctt aaa aat aat cct aca tgatgtaaga taaatatata tttctaaaat 3770  
 Leu Lys Asn Asn Pro Thr  
 1200 1205

tatttttttt ttatacttta atgtgtacaa taaaatatat atctaaatat attttatttg 3830

tacgcttttt tttttttttt ttttaattggt atttttgtat at 3872

&lt;210&gt; 4

&lt;211&gt; 1205

&lt;212&gt; PRT

&lt;213&gt; Plasmodium falciparum

&lt;400&gt; 4

Met Ile Phe Asn Tyr Val Phe Phe Lys Asn Phe Val Pro Val Val Leu  
 1 5 10 15

Tyr Ile Leu Leu Ile Ile Tyr Ile Asn Leu Asn Gly Met Asn Asn Lys  
 20 25 30

Asn Gln Ile Lys Thr Glu Lys Ile Tyr Ile Lys Lys Leu Asn Arg Leu  
 35 40 45

Ser Arg Lys Asn Ser Leu Cys Ser Ser Lys Asn Lys Ile Ala Cys Leu  
 50 55 60

Phe Asp Ile Gly Asn Asp Asp Asn Arg Asn Thr Thr Tyr Gly Tyr Asn  
 65 70 75 80

Val Asn Val Lys Asn Asp Asp Ile Asn Ser Leu Leu Lys Asn Asn Tyr  
 85 90 95

Ser Asn Lys Leu Tyr Met Asp Lys Arg Lys Asn Ile Asn Asn Val Ile  
 100 105 110

Ser Thr Asn Lys Ile Ser Gly Ser Ile Ser Asn Ile Cys Ser Arg Asn  
 115 120 125

WO 00/17233

PCT/EP99/07055

- 12 -

Gln Lys Glu Asn Glu Gln Lys Arg Asn Lys Gln Arg Cys Leu Thr Gln  
 130 135 140

Cys His Thr Tyr Asn Met Ser His Glu Gln Asp Lys Leu Ala Asn Asp  
 145 150 155 160

Asn Asn Arg Asn Asn Lys Lys Asn Phe Asn Leu Leu Phe Ile Asn Tyr  
 165 170 175

Phe Asn Leu Lys Arg Met Lys Asn Ser Leu Leu Asn Lys Asp Asn Phe  
 180 185 190

Phe Tyr Cys Lys Glu Lys Lys Leu Ser Phe Leu His Lys Ala Tyr Lys  
 195 200 205

Lys Lys Asn Cys Thr Phe Gln Asn Tyr Ser Leu Lys Arg Lys Ser Asn  
 210 215 220

Arg Asp Ser His Lys Leu Phe Ser Gly Glu Phe Asp Asp Tyr Thr Asn  
 225 230 235 240

Asn Asn Ala Leu Tyr Glu Ser Glu Lys Lys Glu Tyr Ile Thr Leu Asn  
 245 250 255

Asn Asn Asn Lys Asn Asn Asn Asn Lys Asn Asn Asp Asn Lys Asn Asn  
 260 265 270

Asp Asn Asn Asp Tyr Asn Asn Asn Asn Ser Cys Asn Asn Leu Gly Glu  
 275 280 285

Arg Ser Asn His Tyr Asp Asn Tyr Gly Gly Asp Asn Asn Asn Pro Cys  
 290 295 300

Asn Asn Asn Asn Asp Lys Tyr Asp Ile Gly Lys Tyr Phe Lys Gln Ile  
 305 310 315 320

Asn Thr Phe Ile Asn Ile Asp Glu Tyr Lys Thr Ile Tyr Gly Asp Glu  
 325 330 335

Ile Tyr Lys Glu Ile Tyr Glu Leu Tyr Val Glu Arg Asn Ile Pro Glu  
 340 345 350

Tyr Tyr Glu Arg Lys Tyr Phe Ser Glu Asp Ile Lys Lys Ser Val Leu  
 355 360 365

Phe Asp Ile Asp Lys Tyr Asn Asp Val Glu Phe Glu Lys Ala Ile Lys  
 370 375 380

Glu Glu Phe Ile Asn Asn Gly Val Tyr Ile Asn Asn Ile Asp Asn Thr  
 385 390 395 400

Tyr Tyr Lys Lys Glu Asn Ile Leu Ile Met Lys Lys Ile Leu His Tyr  
 405 410 415

Phe Pro Leu Leu Lys Leu Ile Asn Asn Pro Ser Asp Leu Lys Lys Leu  
 420 425 430

WO 00/17233

PCT/EP99/07055

- 13 -

Lys Lys Gln Tyr Leu Pro Leu Leu Ala His Glu Leu Lys Ile Phe Leu  
 435 440 445  
 Phe Phe Ile Val Asn Ile Thr Gly Gly His Phe Ser Ser Val Leu Ser  
 450 455 460  
 Ser Leu Glu Ile Gln Leu Leu Leu Leu Tyr Ile Phe Asn Gln Pro Tyr  
 465 470 475 480  
 Asp Asn Val Ile Tyr Asp Ile Gly His Gln Ala Tyr Val His Lys Ile  
 485 490 495  
 Leu Thr Gly Arg Lys Leu Leu Phe Leu Ser Leu Arg Asn Lys Lys Gly  
 500 505 510  
 Ile Ser Gly Phe Leu Asn Ile Phe Glu Ser Ile Tyr Asp Lys Phe Gly  
 515 520 525  
 Ala Gly His Ser Ser Thr Ser Leu Ser Ala Ile Gln Gly Tyr Tyr Glu  
 530 535 540  
 Ala Glu Trp Gln Val Lys Asn Lys Glu Lys Tyr Gly Asn Gly Asp Ile  
 545 550 555 560  
 Glu Ile Ser Asp Asn Ala Asn Val Thr Asn Asn Glu Arg Ile Phe Gln  
 565 570 575  
 Lys Gly Ile His Asn Asp Asn Asn Ile Asn Asn Asn Ile Asn Asn Asn  
 580 585 590  
 Asn Tyr Ile Asn Pro Ser Asp Val Val Gly Arg Glu Asn Thr Asn Val  
 595 600 605  
 Pro Asn Val Arg Asn Asp Asn His Asn Val Asp Lys Val His Ile Ala  
 610 615 620  
 Ile Ile Gly Asp Gly Gly Leu Thr Gly Gly Met Ala Leu Glu Ala Leu  
 625 630 635 640  
 Asn Tyr Ile Ser Phe Leu Asn Ser Lys Ile Leu Ile Ile Tyr Asn Asp  
 645 650 655  
 Asn Gly Gln Val Ser Leu Pro Thr Asn Ala Val Ser Ile Ser Gly Asn  
 660 665 670  
 Arg Pro Ile Gly Ser Ile Ser Asp His Leu His Tyr Phe Val Ser Asn  
 675 680 685  
 Ile Glu Ala Asn Ala Gly Asp Asn Lys Leu Ser Lys Asn Ala Lys Glu  
 690 695 700  
 Asn Asn Ile Phe Glu Asn Leu Asn Tyr Asp Tyr Ile Gly Val Val Asn  
 705 710 715 720  
 Gly Asn Asn Thr Glu Glu Leu Phe Lys Val Leu Asn Asn Ile Lys Glu  
 725 730 735

WO 00/17233

- 14 -

Asn Lys Leu Lys Arg Ala Thr Val Leu His Val Arg Thr Lys Lys Ser  
 740 745 750

Asn Asp Phe Ile Asn Ser Lys Ser Pro Ile Ser Ile Leu His Ser Ile  
 755 760 765

Lys Lys Asn Glu Ile Phe Pro Phe Asp Thr Thr Ile Leu Asn Gly Asn  
 770 775 780

Ile His Lys Glu Asn Lys Ile Glu Glu Glu Lys Asn Val Ser Ser Ser  
 785 790 795 800

Thr Lys Tyr Asp Val Asn Asn Lys Asn Asn Lys Asn Asn Asp Asn Ser  
 805 810 815

Glu Ile Ile Lys Tyr Glu Asp Met Phe Ser Lys Glu Thr Phe Thr Asp  
 820 825 830

Ile Tyr Thr Asn Glu Met Leu Lys Tyr Leu Lys Lys Asp Arg Asn Ile  
 835 840 845

Ile Phe Leu Ser Pro Ala Met Leu Gly Gly Ser Gly Leu Val Lys Ile  
 850 855 860

Ser Glu Arg Tyr Pro Asn Asn Val Tyr Asp Val Gly Ile Ala Glu Gln  
 865 870 875 880

His Ser Val Thr Phe Ala Ala Ala Met Ala Met Asn Lys Lys Leu Lys  
 885 890 895

Ile Gln Leu Cys Ile Tyr Ser Thr Phe Leu Gln Arg Ala Tyr Asp Gln  
 900 905 910

Ile Ile His Asp Leu Asn Leu Gln Asn Ile Pro Leu Lys Val Ile Ile  
 915 920 925

Gly Arg Ser Gly Leu Val Gly Glu Asp Gly Ala Thr His Gln Gly Ile  
 930 935 940

Tyr Asp Leu Ser Tyr Leu Gly Thr Leu Asn Asn Ala Tyr Ile Ile Ser  
 945 950 955 960

Pro Ser Asn Gln Val Asp Leu Lys Arg Ala Leu Arg Phe Ala Tyr Leu  
 965 970 975

Asp Lys Asp His Ser Val Tyr Ile Arg Ile Pro Arg Met Asn Ile Leu  
 980 985 990

Ser Asp Lys Tyr Met Lys Gly Tyr Leu Asn Ile His Met Lys Asn Glu  
 995 1000 1005

Ser Lys Asn Ile Asp Val Asn Val Asp Ile Asn Asp Asp Val Asp Lys  
 1010 1015 1020

Tyr Ser Glu Glu Tyr Met Asp Asp Asp Asn Phe Ile Lys Ser Phe Ile  
 025 1030 1035 1040

WO 00/17233

PCT/EP99/07055

- 15 -

Gly Lys Ser Arg Ile Ile Lys Met Asp Asn Glu Asn Asn Asn Thr Asn  
 1045 1050 1055

Glu His Tyr Ser Ser Arg Gly Asp Thr Gln Thr Lys Lys Lys Lys Val  
 1060 1065 1070

Cys Ile Phe Asn Met Gly Ser Met Leu Phe Asn Val Ile Asn Ala Ile  
 1075 1080 1085

Lys Glu Ile Glu Lys Glu Gln Tyr Ile Ser His Asn Tyr Ser Phe Ser  
 1090 1095 1100

Ile Val Asp Met Ile Phe Leu Asn Pro Leu Asp Lys Asn Met Ile Asp  
 1105 1110 1115 1120

His Val Ile Lys Gln Asn Lys His Gln Tyr Leu Ile Thr Tyr Glu Asp  
 1125 1130 1135

Asn Thr Ile Gly Gly Phe Ser Thr His Phe Asn Asn Tyr Leu Ile Glu  
 1140 1145 1150

Asn Asn Tyr Ile Thr Lys His Asn Leu Tyr Val His Asn Ile Tyr Leu  
 1155 1160 1165

Ser Asn Glu Pro Ile Glu His Ala Ser Phe Lys Asp Gln Gln Glu Val  
 1170 1175 1180

Val Lys Met Asp Lys Cys Ser Leu Val Asn Arg Ile Lys Asn Tyr Leu  
 1185 1190 1195 1200

Lys Asn Asn Pro Thr  
 1205

&lt;210&gt; 5

&lt;211&gt; 3147

&lt;212&gt; DNA

&lt;213&gt; Plasmodium falciparum

&lt;220&gt;

&lt;221&gt; CDS

&lt;222&gt; (199)..(2670)

&lt;400&gt; 5

tttcattttt ctttaccac atatatat atatatat aatatata tataatatta 60

tatatttgat atatgattta aaattgtaac ataaaaaaaa taattatatt aaatatgtgt 120

atacatctcc aacatataaa tattattttt tattattatt ttttttttt tttttcataa 180

tgccgaata accacaaa atg agt tat ata aaa aga ctg att ctt ttt atg 231  
 Met Ser Tyr Ile Lys Arg Leu Ile Leu Phe Met  
 1 5 10

tta ctg ttt tat tct cat gta aaa att aaa aaa tta ttt att aaa att 279  
 Leu Leu Phe Tyr Ser His Val Lys Ile Lys Lys Leu Phe Ile Lys Ile  
 15 20 25

WO 00/17233

PCT/EP99/07055

-16-

tct aat gta aac ata ttt ttt gca gaa gca aag aaa aat gga aaa aag	327
Ser Asn Val Asn Ile Phe Phe Ala Glu Ala Lys Lys Asn Gly Lys Lys	
30 35 40	
gaa ttc ttt ctt ttt tta cta aat ata aaa aaa aat agc caa cag aaa	375
Glu Phe Phe Leu Phe Leu Leu Asn Ile Lys Lys Asn Ser Gln Gln Lys	
45 50 55	
aaa act tat cat att acc aaa agg aat acc ata aat aaa agt gat ttt	423
Lys Thr Tyr His Ile Thr Lys Arg Asn Thr Ile Asn Lys Ser Asp Phe	
60 65 70 75	
tta tat tct tta cta aat gaa gaa ggg aat tct tca aaa aag gaa tat	471
Leu Tyr Ser Leu Leu Asn Glu Glu Gly Asn Ser Ser Lys Lys Glu Tyr	
80 85 90	
aaa aat tta aaa gat gaa gaa aaa tat aat atc ata caa aat ata aaa	519
Lys Asn Leu Lys Asp Glu Glu Lys Tyr Asn Ile Ile Gln Asn Ile Lys	
95 100 105	
aaa tat tgt gaa tgt act aaa aaa tat aaa agg ctc cca aca cga gaa	567
Lys Tyr Cys Glu Cys Thr Lys Lys Tyr Lys Arg Leu Pro Thr Arg Glu	
110 115 120	
gta gtt att gga aat gtt aaa att gga gga aat aat aaa ata gct att	615
Val Val Ile Gly Asn Val Lys Ile Gly Gly Asn Asn Lys Ile Ala Ile	
125 130 135	
caa act atg gct agc tgt gat aca aga aat gta gaa gaa tgt gta tat	663
Gln Thr Met Ala Ser Cys Asp Thr Arg Asn Val Glu Glu Cys Val Tyr	
140 145 150 155	
caa att aga aaa tgt aaa gat ttg ggt gct gac att gta agg ttg act	711
Gln Ile Arg Lys Cys Lys Asp Leu Gly Ala Asp Ile Val Arg Leu Thr	
160 165 170	
gtt caa gga gtt caa gaa gca caa gct agt tat cat att aaa gaa aaa	759
Val Gln Gly Val Gln Glu Ala Gln Ala Ser Tyr His Ile Lys Glu Lys	
175 180 185	
tta tta tct gaa aat gta aat atc cca tta gta gca gat att cat ttt	807
Leu Leu Ser Glu Asn Val Asn Ile Pro Leu Val Ala Asp Ile His Phe	
190 195 200	
aat cct aaa ata gct tta atg gca gct gat gtg ttt gaa aaa att cga	855
Asn Pro Lys Ile Ala Leu Met Ala Ala Asp Val Phe Glu Lys Ile Arg	
205 210 215	
gtg aat cca gga aat tat gtt gat gga aga aaa aaa tgg ata gat aaa	903
Val Asn Pro Gly Asn Tyr Val Asp Gly Arg Lys Lys Trp Ile Asp Lys	
220 225 230 235	
gtt tat aaa aat aaa gaa gaa ttt gat gaa ggg aaa tta ttt ata aaa	951
Val Tyr Lys Thr Lys Glu Glu Phe Asp Glu Gly Lys Leu Phe Ile Lys	
240 245 250	

WO 00/17233

PCT/EP99/07055

- 17 -

gaa aaa ttt gta cca tta att gaa aaa tgt aaa aga tta aat aga gca	999
Glu Lys Phe Val Pro Leu Ile Glu Lys Cys Lys Arg Leu Asn Arg Ala	
255 260 265	
ata aga att gga aca aat cat gga tcc ctt tca tct cga gta tta tca	1047
Ile Arg Ile Gly Thr Asn His Gly Ser Leu Ser Ser Arg Val Leu Ser	
270 275 280	
tat tat gga gat aca cca tta ggt atg gta gaa tct gct ttt gag ttt	1095
Tyr Tyr Gly Asp Thr Pro Leu Gly Met Val Glu Ser Ala Phe Glu Phe	
285 290 295	
tct gat tta tgt att gaa aac aat ttt tac aat ctt gtt ttt tct atg	1143
Ser Asp Leu Cys Ile Glu Asn Asn Phe Tyr Asn Leu Val Phe Ser Met	
300 305 310 315	
aaa gct tct aat gct tat gtt atg ata caa tct tat aga tta tta gta	1191
Lys Ala Ser Asn Ala Tyr Val Met Ile Gln Ser Tyr Arg Leu Leu Val	
320 325 330	
tct aaa caa tat gaa aga aat atg atg ttc cct ata cat tta gga gtt	1239
Ser Lys Gln Tyr Glu Arg Asn Met Met Phe Pro Ile His Leu Gly Val	
335 340 345	
aca gaa gca gga ttt ggt gat aat gga aga ata aaa tct tat tta ggt	1287
Thr Glu Ala Gly Phe Gly Asp Asn Gly Arg Ile Lys Ser Tyr Leu Gly	
350 355 360	
ata gga tct tta tta tat gat ggt ata gga gat acc att cgt ata tcc	1335
Ile Gly Ser Leu Leu Tyr Asp Gly Ile Gly Asp Thr Ile Arg Ile Ser	
365 370 375	
tta aca gaa gat cct tgg gaa gag tta act cct tgt aaa aaa tta gtt	1383
Leu Thr Glu Asp Pro Trp Glu Glu Leu Thr Pro Cys Lys Lys Leu Val	
380 385 390 395	
gaa aat tta aag aaa aga ata ttt tat aat gaa aat ttt aaa gaa gat	1431
Glu Asn Leu Lys Lys Arg Ile Phe Tyr Asn Glu Asn Phe Lys Glu Asp	
400 405 410	
aat gaa tta aaa aat aat gaa atg gat acc aaa aat cta tta aat ttt	1479
Asn Glu Leu Lys Asn Asn Glu Met Asp Thr Lys Asn Leu Leu Asn Phe	
415 420 425	
gaa gaa aat tat cga aat ttt aat aat ata aaa aaa aga aat gta gaa	1527
Glu Glu Asn Tyr Arg Asn Phe Asn Asn Ile Lys Lys Arg Asn Val Glu	
430 435 440	
aaa aat aat aat gta tta cat gaa gag tgc act ata ggt aat gta gta	1575
Lys Asn Asn Asn Val Leu His Glu Glu Cys Thr Ile Gly Asn Val Val	
445 450 455	
acc ata aaa gag tta gaa gat tct ctg caa att ttt aaa gat tta aat	1623
Thr Ile Lys Glu Leu Glu Asp Ser Leu Gln Ile Phe Lys Asp Leu Asn	
460 465 470 475	



WO 00/17233

PCT/EP99/07055

- 18 -

tta gaa gta gat tca aat gga aat ttg aaa aag gga gcc aaa aca act	1671
Leu Glu Val Asp Ser Asn Gly Asn Leu Lys Lys Gly Ala Lys Thr Thr	
480 485 490	
gat atg gtt att ata aat gat ttt cat aat ata aca aat tta gga aaa	1719
Asp Met Val Ile Ile Asn Asp Phe His Asn Ile Thr Asn Leu Gly Lys	
495 500 505	
aaa act gtg gat aaa tta atg caa gtg gga att aat ata gta gtt caa	1767
Lys Thr Val Asp Lys Leu Met Gln Val Gly Ile Asn Ile Val Val Gln	
510 515 520	
tat gaa cca cat aat ata gaa ttt ata gaa aaa atg gaa cca aat aat	1815
Tyr Glu Pro His Asn Ile Glu Phe Ile Glu Lys Met Glu Pro Asn Asn	
525 530 535	
gat aat aat aat aat aat aat aat aat aat ata tta ttt tat gtg gat	1863
Asp Asn Asn Asn Asn Asn Asn Asn Asn Ile Leu Phe Tyr Val Asp	
540 545 550 555	
ata aaa aat att atg aac agt tca gaa aaa aat att aaa tta agt aat	1911
Ile Lys Asn Ile Met Asn Ser Ser Glu Lys Asn Ile Lys Leu Ser Asn	
560 565 570	
tct aaa gga tat gga tta att tta aac gga aaa gaa gat ata caa acc	1959
Ser Lys Gly Tyr Gly Leu Ile Leu Asn Gly Lys Glu Asp Ile Gln Thr	
575 580 585	
ata aaa aaa ata aaa gaa tta aat cgt cgt cct tta ttc att cta tta	2007
Ile Lys Lys Ile Lys Glu Leu Asn Arg Arg Pro Leu Phe Ile Leu Leu	
590 595 600	
aaa tca gat aac ata tat gaa cat gta tta ata acc aga aga att aat	2055
Lys Ser Asp Asn Ile Tyr Glu His Val Leu Ile Thr Arg Arg Ile Asn	
605 610 615	
gaa ctt tta caa tcc tta aat ata aat ata cct tat ata cat tat gtt	2103
Glu Leu Leu Gln Ser Leu Asn Ile Asn Ile Pro Tyr Ile His Tyr Val	
620 625 630 635	
gat att aat tca aac aat tat gat gat ata tta gtt aat tca aca tta	2151
Asp Ile Asn Ser Asn Asn Tyr Asp Asp Ile Leu Val Asn Ser Thr Leu	
640 645 650	
tat gca gga agt tgt ttg atg gat tta atg ggg gat ggt ctt att gtt	2199
Tyr Ala Gly Ser Cys Leu Met Asp Leu Met Gly Asp Gly Leu Ile Val	
655 660 665	
aac gta act aat gat gtt ctt aca aat aaa aaa aag ata gaa aca aaa	2247
Asn Val Thr Asn Asp Val Leu Thr Asn Lys Lys Lys Ile Glu Thr Lys	
670 675 680	
tat gat gaa aaa gaa gaa gta gag gaa gag gga aac aat aaa gat att	2295
Tyr Asp Glu Lys Glu Glu Val Glu Glu Glu Gly Asn Asn Lys Asp Ile	
685 690 695	



WO 00/17233

PCT/EP99/07055

- 19 -

cat aga ctt ttg agc aga gtt gca tta aat tca ttt tta aca tta aat 2343  
 His Arg Leu Leu Ser Arg Val Ala Leu Asn Ser Phe Leu Thr Leu Asn  
 700 705 710 715  
 att tta caa gat aca aga ata cgt tta ttt aaa aca gat tat ata gcc 2391  
 Ile Leu Gln Asp Thr Arg Ile Arg Leu Phe Lys Thr Asp Tyr Ile Ala  
 720 725 730  
 tgc cca tct tgt gga aga act tta ttt aat ata caa gaa act act aaa 2439  
 Cys Pro Ser Cys Gly Arg Thr Leu Phe Asn Ile Gln Glu Thr Thr Lys  
 735 740 745  
 aaa att atg aaa tta aca ggg cac tta aaa ggc gtt aaa att gca gtc 2487  
 Lys Ile Met Lys Leu Thr Gly His Leu Lys Gly Val Lys Ile Ala Val  
 750 755 760  
 atg gga tgt att gtt aat ggt ata gga gaa atg gca gat gca cat ttt 2535  
 Met Gly Cys Ile Val Asn Gly Ile Gly Glu Met Ala Asp Ala His Phe  
 765 770 775  
 ggt tat gtt ggt agt gca cct aaa aaa att gat tta tat tat ggt aaa 2583  
 Gly Tyr Val Gly Ser Ala Pro Lys Lys Ile Asp Leu Tyr Tyr Gly Lys  
 780 785 790 795  
 gag tta gta gaa aga aat ata cct gag gaa gaa gct tgt gat aaa ttg 2631  
 Glu Leu Val Glu Arg Asn Ile Pro Glu Glu Glu Ala Cys Asp Lys Leu  
 800 805 810  
 ata gaa tta att aaa aaa cat aac aaa tgg aaa gat cca taaattgaat 2680  
 Ile Glu Leu Ile Lys Lys His Asn Lys Trp Lys Asp Pro  
 815 820  
 atggacaagt atttatattat ttatttatct tatatataat atattataaa tttttc gatg 2740  
 tatttttccct tttaaaattt tatttttttt ttattttttt ttttgaagta atatttataa 2800  
 tgcatacata atattaaaat gtgtattata taataatatac attttattgt tatttttaaaa 2860  
 gactaatacc aagaacaatt ttttaataat cattcctata acttggttaa tatatatata 2920  
 tatatatata tatttattta tttatattta tatttattta tttttggtat atgaaaagta 2980  
 aaaatataat aatttaaaag tatttacaaa ataaataata ttatatatct gtttttatat 3040  
 atatgttaat ggaaaaggag aaaataaata aataaaacaa acaaaataac atatatatat 3100  
 atatatatat actgaatgag aaagaaaaaa aaaagaaaag gatacga 3147

&lt;210&gt; 6

&lt;211&gt; 824

&lt;212&gt; PRT

&lt;213&gt; Plasmodium falciparum

&lt;400&gt; 6

Met Ser Tyr Ile Lys Arg Leu Ile Leu Phe Met Leu Leu Phe Tyr Ser  
 1 5 10 15

WO 00/17233

PCT/EP99/07055

- 20 -

His Val Lys Ile Lys Lys Leu Phe Ile Lys Ile Ser Asn Val Asn Ile  
 20 25 30  
 Phe Phe Ala Glu Ala Lys Lys Asn Gly Lys Lys Glu Phe Phe Leu Phe  
 35 40 45  
 Leu Leu Asn Ile Lys Lys Asn Ser Gln Gln Lys Lys Thr Tyr His Ile  
 50 55 60  
 Thr Lys Arg Asn Thr Ile Asn Lys Ser Asp Phe Leu Tyr Ser Leu Leu  
 65 70 75 80  
 Asn Glu Glu Gly Asn Ser Ser Lys Lys Glu Tyr Lys Asn Leu Lys Asp  
 85 90 95  
 Glu Glu Lys Tyr Asn Ile Ile Gln Asn Ile Lys Lys Tyr Cys Glu Cys  
 100 105 110  
 Thr Lys Lys Tyr Lys Arg Leu Pro Thr Arg Glu Val Val Ile Gly Asn  
 115 120 125  
 Val Lys Ile Gly Gly Asn Asn Lys Ile Ala Ile Gln Thr Met Ala Ser  
 130 135 140  
 Cys Asp Thr Arg Asn Val Glu Glu Cys Val Tyr Gln Ile Arg Lys Cys  
 145 150 155 160  
 Lys Asp Leu Gly Ala Asp Ile Val Arg Leu Thr Val Gln Gly Val Gln  
 165 170 175  
 Glu Ala Gln Ala Ser Tyr His Ile Lys Glu Lys Leu Leu Ser Glu Asn  
 180 185 190  
 Val Asn Ile Pro Leu Val Ala Asp Ile His Phe Asn Pro Lys Ile Ala  
 195 200 205  
 Leu Met Ala Ala Asp Val Phe Glu Lys Ile Arg Val Asn Pro Gly Asn  
 210 215 220  
 Tyr Val Asp Gly Arg Lys Lys Trp Ile Asp Lys Val Tyr Lys Thr Lys  
 225 230 235 240  
 Glu Glu Phe Asp Glu Gly Lys Leu Phe Ile Lys Glu Lys Phe Val Pro  
 245 250 255  
 Leu Ile Glu Lys Cys Lys Arg Leu Asn Arg Ala Ile Arg Ile Gly Thr  
 260 265 270  
 Asn His Gly Ser Leu Ser Ser Arg Val Leu Ser Tyr Tyr Gly Asp Thr  
 275 280 285  
 Pro Leu Gly Met Val Glu Ser Ala Phe Glu Phe Ser Asp Leu Cys Ile  
 290 295 300  
 Glu Asn Asn Phe Tyr Asn Leu Val Phe Ser Met Lys Ala Ser Asn Ala  
 305 310 315 320  
 Tyr Val Met Ile Gln Ser Tyr Arg Leu Leu Val Ser Lys Gln Tyr Glu

325	330	335
Arg Asn Met Met Phe Pro Ile His Leu Gly Val Thr Glu Ala Gly Phe		
340	345	350
Gly Asp Asn Gly Arg Ile Lys Ser Tyr Leu Gly Ile Gly Ser Leu Leu		
355	360	365
Tyr Asp Gly Ile Gly Asp Thr Ile Arg Ile Ser Leu Thr Glu Asp Pro		
370	375	380
Trp Glu Glu Leu Thr Pro Cys Lys Lys Leu Val Glu Asn Leu Lys Lys		
385	390	395
Arg Ile Phe Tyr Asn Glu Asn Phe Lys Glu Asp Asn Glu Leu Lys Asn		
405	410	415
Asn Glu Met Asp Thr Lys Asn Leu Leu Asn Phe Glu Glu Asn Tyr Arg		
420	425	430
Asn Phe Asn Asn Ile Lys Lys Arg Asn Val Glu Lys Asn Asn Asn Val		
435	440	445
Leu His Glu Glu Cys Thr Ile Gly Asn Val Val Thr Ile Lys Glu Leu		
450	455	460
Glu Asp Ser Leu Gln Ile Phe Lys Asp Leu Asn Leu Glu Val Asp Ser		
465	470	475
Asn Gly Asn Leu Lys Lys Gly Ala Lys Thr Thr Asp Met Val Ile Ile		
485	490	495
Asn Asp Phe His Asn Ile Thr Asn Leu Gly Lys Lys Thr Val Asp Lys		
500	505	510
Leu Met Gln Val Gly Ile Asn Ile Val Val Gln Tyr Glu Pro His Asn		
515	520	525
Ile Glu Phe Ile Glu Lys Met Glu Pro Asn Asn Asp Asn Asn Asn Asn		
530	535	540
Asn Asn Asn Asn Asn Ile Leu Phe Tyr Val Asp Ile Lys Asn Ile Met		
545	550	555
Asn Ser Ser Glu Lys Asn Ile Lys Leu Ser Asn Ser Lys Gly Tyr Gly		
565	570	575
Leu Ile Leu Asn Gly Lys Glu Asp Ile Gln Thr Ile Lys Lys Ile Lys		
580	585	590
Glu Leu Asn Arg Arg Pro Leu Phe Ile Leu Leu Lys Ser Asp Asn Ile		
595	600	605
Tyr Glu His Val Leu Ile Thr Arg Arg Ile Asn Glu Leu Leu Gln Ser		
610	615	620
Leu Asn Ile Asn Ile Pro Tyr Ile His Tyr Val Asp Ile Asn Ser Asn		
625	630	635
		640

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PCT/EP99/07055

- 22 -

Asn Tyr Asp Asp Ile Leu Val Asn Ser Thr Leu Tyr Ala Gly Ser Cys  
 645 650 655

Leu Met Asp Leu Met Gly Asp Gly Leu Ile Val Asn Val Thr Asn Asp  
 660 665 670

Val Leu Thr Asn Lys Lys Lys Ile Glu Thr Lys Tyr Asp Glu Lys Glu  
 675 680 685

Glu Val Glu Glu Glu Gly Asn Asn Lys Asp Ile His Arg Leu Leu Ser  
 690 695 700

Arg Val Ala Leu Asn Ser Phe Leu Thr Leu Asn Ile Leu Gln Asp Thr  
 705 710 715 720

Arg Ile Arg Leu Phe Lys Thr Asp Tyr Ile Ala Cys Pro Ser Cys Gly  
 725 730 735

Arg Thr Leu Phe Asn Ile Gln Glu Thr Thr Lys Lys Ile Met Lys Leu  
 740 745 750

Thr Gly His Leu Lys Gly Val Lys Ile Ala Val Met Gly Cys Ile Val  
 755 760 765

Asn Gly Ile Gly Glu Met Ala Asp Ala His Phe Gly Tyr Val Gly Ser  
 770 775 780

Ala Pro Lys Lys Ile Asp Leu Tyr Tyr Gly Lys Glu Leu Val Glu Arg  
 785 790 795 800

Asn Ile Pro Glu Glu Glu Ala Cys Asp Lys Leu Ile Glu Leu Ile Lys  
 805 810 815

Lys His Asn Lys Trp Lys Asp Pro  
 820